UNITED STATES DEPARTMENT OF COMMERCE United States Patent and Trademark Office Address: COMMISSIONER FOR PATENTS P.O. Box 1450 Alexandria, Virginia 22313-1450 www.uspto.gov

APPLICATION NO.	FILING DATE	FIRST NAMED INVENTOR	ATTORNEY DOCKET NO.	CONFIRMATION NO.
10/578,673	05/09/2006	Amr Ali Al-Hossary		3883
Ali Mokhtar Al Hossary 12 El- Morsy Street ard Eel-Etr Al Sananeyyah Domietta post code, 34713 EGYPT			EXAMINER	
			PORTNER, VIRGINIA ALLEN	
			ART UNIT	PAPER NUMBER
			1645	
			MAIL DATE	DELIVERY MODE
			06/18/2009	PAPER

Please find below and/or attached an Office communication concerning this application or proceeding.

The time period for reply, if any, is set in the attached communication.

	Application No.	Applicant(s)
	10/578,673	AL-HOSSARY, AMR ALI
Office Action Summary	Examiner	Art Unit
	GINNY PORTNER	1645
The MAILING DATE of this communication appeariod for Reply	ppears on the cover sheet with the o	correspondence address
A SHORTENED STATUTORY PERIOD FOR REP WHICHEVER IS LONGER, FROM THE MAILING - Extensions of time may be available under the provisions of 37 CFR 1 after SIX (6) MONTHS from the mailing date of this communication. - If NO period for reply is specified above, the maximum statutory perio - Failure to reply within the set or extended period for reply will, by statu. Any reply received by the Office later than three months after the mail earned patent term adjustment. See 37 CFR 1.704(b).	DATE OF THIS COMMUNICATION 1.136(a). In no event, however, may a reply be tind d will apply and will expire SIX (6) MONTHS from the cause the application to become ABANDONE	N. mely filed the mailing date of this communication. ED (35 U.S.C. § 133).
Status		
1) ☐ Responsive to communication(s) filed on <u>01</u> 2a) ☐ This action is FINAL . 2b) ☐ Th 3) ☐ Since this application is in condition for allow closed in accordance with the practice under	nis action is non-final. vance except for formal matters, pro	
Disposition of Claims		
4) Claim(s) 19-27 is/are pending in the application 4a) Of the above claim(s) is/are withdrest spling is/are allowed. 5) Claim(s) is/are allowed. 6) Claim(s) 19-27 is/are rejected. 7) Claim(s) is/are objected to. 8) Claim(s) are subject to restriction and are subject to restriction and are subject to restriction.	rawn from consideration. /or election requirement.	
9) The specification is objected to by the Examir 10) The drawing(s) filed on is/are: a) according a control and applicant may not request that any objection to the Replacement drawing sheet(s) including the correct should be control at the control and the correct should be control and the correct should be control at the control and the con	ccepted or b) objected to by the e drawing(s) be held in abeyance. Se ection is required if the drawing(s) is ob	e 37 CFR 1.85(a). ejected to. See 37 CFR 1.121(d).
Priority under 35 U.S.C. § 119		
12) ☐ Acknowledgment is made of a claim for foreign a) ☐ All b) ☐ Some * c) ☐ None of: 1. ☐ Certified copies of the priority document a. ☐ Certified copies of the priority document a. ☐ Copies of the certified copies of the priority document application from the International Bure * See the attached detailed Office action for a list	nts have been received. nts have been received in Applicat iority documents have been receive au (PCT Rule 17.2(a)).	ion No ed in this National Stage
Attachment(s) 1) Notice of References Cited (PTO-892) 2) Notice of Draftsperson's Patent Drawing Review (PTO-948) 3) Information Disclosure Statement(s) (PTO/SB/08) Paper No(s)/Mail Date	4) Interview Summary Paper No(s)/Mail D 5) Notice of Informal F 6) Other:	ate

Application/Control Number: 10/578,673 Page 2

Art Unit: 1645

DETAILED ACTION

Claims 19-27 are pending.

Continued Examination Under 37 CFR 1.114

1. A request for continued examination under 37 CFR 1.114, including the fee set forth in

37 CFR 1.17(e), was filed in this application after final rejection. Since this application is

eligible for continued examination under 37 CFR 1.114, and the fee set forth in 37 CFR 1.17(e)

has been timely paid, the finality of the previous Office action has been withdrawn pursuant to

37 CFR 1.114. Applicant's submission filed on June 1, 2009 has been entered.

Rejections Maintained

2. Claims 10,12-16 and 18 are rejected under 35 U.S.C. 112, second paragraph, as being

indefinite for failing to particularly point out and distinctly claim the subject matter which

applicant regards as the invention, for reasons of record and in light of the fact that the prior art

filter has all of the claimed structural elements.

3. Claims 10-18 are rejected under 35 U.S.C. 102(b) as being anticipated by Cardone et al,

US Pat. 5,573,957, for reasons of record.

Please Note: The instant Application has been assigned to a new examiner, that has considered

the prior office action, the pending claims, the instant Specification, and the prior art of record.

The instant examiner will be proposing claims that will define over the prior art of record, and

could be allowable if submitted in response to this office action.

Proposed Claim Amendments: Cancel pending claims 19-27.

Art Unit: 1645

Claim 28 (New) An intravenous device comprising a rod of a shape and size for insertion into a vein, the rod being coated with a sheath, and the sheath being coated with anti-myoglobin antibodies able to remove myoglobin from circulating blood.

Claim 29. (New) The intravenous device of claim 28, wherein the rod is a wire.

Claim 30. (New) The intravenous device of claim 28, wherein the sheath is latex.

Claim 31. (New) The intravenous device of claim 28, wherein the anti-myoglobin antibodies are latex particle immobilized anti-myoglobin antibodies.

Claim 33. (New) A cannula comprising the intravenous device of claim 1.

Claim 34. (New) The cannula of claim 33, wherein the cannula comprises a tip.

Claim 35: (New) The cannula of claim 34, wherein the cannula is a cylindrical shape with a central core into which the intravenous device is inserted, wherein the intravenous device can protrude from the tip end of the cannula.

Claim 36. (New) The cannula of claim 35, wherein the anti-myoglobin antibodies are coated over the whole surface area that protrudes from the tip end of the cannula.

Claim 37: (New) A method comprising the steps of:

Percutaneously introducing the cannula of claim 33 into a vein of a patient,

Removing the intravenous device (see page 8, bottom two paragraphs) from the patient after a period of time during which myoglobin is trapped by the anti-myoglobin antibodies.

Optionally, if needed, reintroducing another intravenous device through the central core of cannula into the patient's vein to trap additional circulating myoglobin, then removing the cannula and intravenous device from the patient.

Art Unit: 1645

Claim 38: (New) The method of claim 37, wherein the patient is a patient at risk of ARF (what does this mean?), rhabdomyolysis, or myogloinemia.

Page 4

Claim 39: (New) The method of claim 38, wherein the risk of ARF is associated with earth quakes, orthopedic problems, alcohol and heroin use, occlusion of muscular vessels, crush injuries, compartment syndrome or reperfusion injury in a limb.

Claim 40. (New) The method of claim 39, wherein the reperfusion injury in a limb is associated with operative reperfusion.

Claim 41: (New) The method of claim 37, wherein the vein is the vinae cava, internal jugular vein or femoral vein.

Specification

- 4. The disclosure is objected to because of the following informalities:
- 5. The instant Specification recites various abbreviations, the meanings of which are unclear. Please provide the full meaning/definition of the following terms:
 - i. IV (title)
 - ii. I.S.F. (page 1)
 - iii. mmHg (page 2)
 - iv. A.R.F. (page 2)
 - v. RBF (page 4)
 - vi. GFR (page 4)
 - vii. CC (page 6)
 - viii. I.J.V. (page 8)
 - ix. I.V.C. (page 8)
 - x. D.V.T. (page 8)
 - xi. S.V.C (page 8)

Appropriate correction is required.

Conclusion

6. The prior art made of record and not relied upon is considered pertinent to applicant's disclosure. Various references are being made of record to show microdialysis filters, and intravenous cannulas.

Any inquiry concerning this communication or earlier communications from the examiner should be directed to GINNY PORTNER whose telephone number is (571)272-0862. The examiner can normally be reached on flextime, but usually M-F, alternate Fridays off.

If attempts to reach the examiner by telephone are unsuccessful, the examiner's supervisor, Robert Mondesi can be reached on 571-272-0956. The fax phone number for the organization where this application or proceeding is assigned is 571-273-8300.

Information regarding the status of an application may be obtained from the Patent Application Information Retrieval (PAIR) system. Status information for published applications may be obtained from either Private PAIR or Public PAIR. Status information for unpublished applications is available through Private PAIR only. For more information about the PAIR system, see http://pair-direct.uspto.gov. Should you have questions on access to the Private PAIR system, contact the Electronic Business Center (EBC) at 866-217-9197 (toll-free). If you would like assistance from a USPTO Customer Service Representative or access to the automated information system, call 800-786-9199 (IN USA OR CANADA) or 571-272-1000.

/Robert B Mondesi/ Supervisory Patent Examiner, Art Unit 1645

/Ginny Portner/ Examiner, Art Unit 1645 June 17, 2009 Application/Control Number: 10/578,673

Page 6

Art Unit: 1645